



Faculty of Manufacturing Engineering

**PREPARATION & CHARACTERIZATION OF
ELECTROPHORETICALLY DEPOSITED BN FILM FOR
SEMICONDUCTOR PACKAGE**

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Master of Science in Manufacturing Engineering

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**PREPARATION & CHARACTERIZATION OF ELECTROPHORETICALLY
DEPOSITED BN FILM FOR SEMICONDUCTOR PACKAGE**

JAYAGANASAN A/L NARAYANASAMY

**A thesis submitted
in fulfillment of the requirements for the degree of Master of Science
in Manufacturing Engineering**

Faculty of Manufacturing Engineering

UNIVERSITI TEKNIKAL MALAYSIA MELAKA

2017

DECLARATION

I declare that this thesis entitled “Preparation & Characterization of Electrophoretically Deposited BN Film for Semiconductor Package” is the result of my own research except as cited in the references. The thesis has not been accepted for any degree and is not concurrently submitted in candidature of any other degree.

Signature :

Name :

Date :

APPROVAL

I hereby declare that I have read this thesis and in my opinion this thesis is sufficient in terms of scope and quality for the award of Master of Science in Manufacturing Engineering.

Signature :

Supervisor Name :

Date :

DEDICATION

To my beloved family, lecturers and friends.

ABSTRACT

Boron Nitride (BN) is used in various applications such as in lubrication, as a releasing agent and also as a thermosetting insulator material and thermal enhancer because of its advanced material properties. Electrophoretic deposition (EPD) method is a new method for semiconductor industries and it has grown in the interest on making use of BN as a thermal interface material and electrical isolation in Transistor Outline (TO) packages. Hexagonal Boron Nitride (h-BN) stability in EPD suspension is essential to produce repeatability and reproducibility in the result of the deposition. However, h-BN particle has less functional group on its side wall to create bonding with the polymer matrix. In order to increase functional groups on its side wall, NaOH surface treatment which has been established by other researchers was performed. The purpose of this study is to prepare and characterise of the h-BN particles in EPD suspension using different types of suspension mediums and binders. The particle size characterization and Field-Emission Scanning Electron Microscopy (FESEM) on the as-received h-BN particle indicates that particle sizes were less than 1 μm but are in agglomerated forms in the De-ionized (DI) water suspension. Sedimentation test method of h-BN particles in four dispersion media (deionized water, Acetic acid solution, Sulphamic Acid & Ammonia) and using different binder (Polyethylene Glycol (PEG), Silane Coupling Agent, Polycationic 1 (PC 1), Polycationic 2 (PC 2). The result showed a combination of deionized water and PC 2 produced the highest stability for h-BN dispersion. Sedimentation test and zeta potential method were used to determine the optimum concentration of PC 2 addition in a h-BN suspension. EPD of h-BN was performed on TO package using different levels of PC 2 concentration (i.e. 0.2 – 1.0 wt%). Characterization of the EPD coating were performed in terms of thickness, microstructure analysis on surface and micrograph from FESEM, and surface roughness. The optimum concentration of PC 2 in order to achieve the highest h-BN stability was in the range of 0.3 – 0.4 wt%, with a corresponding deposition thickness of 8 μm . The obtained thickness was the highest among other samples, and had surface roughness of 570 nm. Critical factors that affected the deposition for h-BN EPD process were suspension ionic conductivity and excess PC 2 concentration. High conductivity and excess PC 2 concentration caused electric double layer of h-BN particles to be compressed thus resulting in a low deposition yield. Therefore, it is recommended that future works use ultra-pure DI water and excess binder of h-BN suspension need to be removed by centrifugal washing before undergoes EPD to reduce conductivity of h-BN suspension. Besides it also helps to achieve a high deposition thickness of h-BN for thermal conductive and electrical isolation application.

ABSTRAK

Boron Nitride (BN) digunakan dalam pelbagai aplikasi seperti dalam pelinciran, sebagai ejen pelepasan dan juga sebagai bahan penebat haba serta pengkonduksian haba kerana sifat-sifat bahan termajunya. Kaedah penyaduran electrophoretik (EPD) adalah satu kaedah baru bagi industri semikonduktor, maka terdapat peningkatan minat ke atas penggunaan BN sebagai bahan antaramuka haba dan penebatan elektrik dalam pakej TO. Kestabilan heksagon Boron Nitride (h-BN) dalam ampaian EPD adalah penting untuk membolehkan pengulangan dan penghasilan semula data penyaduran. Walau bagaimanapun, zarah h-BN mempunyai sedikit kumpulan berfungsi pada permukaan sisi untuk mewujudkan ikatan dengan matrik polimer. Dalam usaha untuk meningkatkan kumpulan berfungsi pada dinding sisi, rawatan permukaan NaOH telah dilakukan sepertimana yang telah diwujudkan oleh penyelidik lain. Tujuan kajian ini adalah untuk menyedia dan mencari zarah h-BN dalam ampaian EPD menggunakan jenis media ampaian dan pengikat berlainan. Pencirian saiz zarah dan Pengimbas Elektron Mikroskop (FESEM) ke atas zarah h-BN mentah menunjukkan saiznya kurang daripada 1 mikron tetapi dalam bentuk gumpalan di dalam ampaian air ternyahion (DI). Kaedah ujian pemendapan zarah h-BN dalam media ampaian berbeza (air ternyahion, pelarut asetik asid, acid sulfamic & Ammonia) dan menggunakan pengikat yang berbeza (poli etilena glikol (PEG), ejen Silane gandingan, Poli kationik 1 (PC 1), Poli kationik 2 (PC 2)) menunjukkan kombinasi air ternyahion dan PC 2 menghasilkan kestabilan paling tinggi berbanding dengan kombinasi medium ampaian dan pengikat lain bagi h-BN. Ujian pemendapan dan kaedah keupayaan zeta telah digunakan untuk menentukan kepekatan PC 2 optimum perlu ditambahkan ke dalam h-BN ampaian. EPD daripada h-BN telah dilakukan ke atas pakej menggunakan tahap kepekatan PC2 berbeza (i.e. 0.2-1.0 wt%). Pencirian saduran EPD termasuk ketebalan, analisis mikrostruktur pada permukaan dan keratan rentas menggunakan FESEM, dan kekasaran permukaan. Kepekatan optimum diperlukan oleh PC 2 untuk mencapai kestabilan tertinggi ialah dalam julat 0.3-0.4 wt%, dengan ketebalan penyaduran berkenaan pada 8 μm . Ketebalan penyaduran yang tertinggi mempunyai kekasaran permukaan sebanyak 570 nm. Faktor kritikal yang memberi kesan kepada penyaduran h-BN EPD adalah kekonduksian ionik dan kepekatan PC 2 berlebihan di dalam ampaian. Faktor ini menyebabkan dwi-lapisan elektrik zarah h-BN dimampatkan, mengakibatkan hasil penyaduran yang rendah. Maka, penyelidikan akan datang disyorkan menggunakan air DI ultra tulen, serta agen pengikat berlebihan bagi ampaian h-BN perlu diemparkan sebelum proses EPD. Kaedah ini akan menyumbang kekonduksian yang rendah, serta membantu mencapai ketebalan penyaduran h-BN yang tinggi untuk aplikasi konduksi haba dan penebatan elektrik.

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**LIST OF ABBREVIATIONS,
SYMBOLS AND NOMENCLATURE**

| | | |
|-------------------------|---|------------------------------|
| μ | - | Electrophoretic Mobility |
| μL | - | Micro Litre |
| μm | - | Micro Meter |
| $\mu\text{S/cm}$ | - | Conductivity Unit |
| A | - | Ampere |
| A | - | Surface Area |
| Al | - | Aluminium |
| Al_2O_3 | - | Aluminum Oxide / Alumina |
| AlN | - | Aluminium Nitride |
| amu | - | Average Molecular Weight |
| B | - | Boron |
| B_2O_3 | - | Boron Trioxide |
| BaTiO_3 | - | Barium Titanium Oxide |
| BCP | - | Biphasic Calcium Phosphate |
| BD | - | Breakdown |
| BDV | - | Breakdown Voltage |
| BJT | - | Bipolar Junction Transistors |
| BN | - | Boron Nitride |

| | | |
|--------------------------------|---|---|
| C | - | Carbon |
| C | - | Celcius |
| C/W | - | Temperature Resistance Unit |
| CaO | - | Calcium Oxide |
| c-BN | - | Cubic Boron Nitride |
| CH ₃ | - | Methyl Group |
| CHO | - | Aldehyde group |
| Cl | - | Chlorine |
| cm ² | - | Centi Meter Square |
| cm ⁻³ | - | Centi Meter Cubic |
| CN | - | Cyanide Functional Group |
| DB | - | Dielectric Breakdown |
| DI | - | Deionized Water |
| E | - | Electric Field Strength |
| EC | - | Electrical Conductivity |
| EDX | - | Energy-dispersive X-ray spectroscopy |
| ELD | - | Electrolytic Deposition |
| EPD | - | Electrophoretic Disposition |
| eV | - | Electron Volt |
| f(Ka) | - | Henry's Function |
| Fe ₂ O ₃ | - | Ferum Oxide |
| FET | - | Field-effect Transistors |
| FTIR | - | Fourier Transform Infrared Spectroscopy |
| g | - | gramme |

| | | |
|--------------------|---|---|
| gm/cm ³ | - | Density unit |
| GPa | - | Giga Pascal |
| H ⁺ | - | Hydrogen Ions |
| H ₂ O | - | Water |
| HA | - | hydroxyapatite |
| h-BN | - | Hexagonal Boron Nitride |
| HCl | - | Hydrochloric Acid |
| HNO ₃ | - | Nitric Acid |
| IC | - | Integrated Circuit |
| IEP | - | Isoelectric Point |
| IGBT | - | Insulated Gate Bipolar Transistor |
| IhBN | - | Isocyanate – Hexagonal BN |
| K | - | Kelvin |
| kHz | - | Kilo Hertz |
| kV | - | Kilo Voltage |
| kV/mm | - | Dielectric Strength Unit |
| kV/s | - | Kilo Voltage per Second (Voltage Breakdown) |
| M | - | Molarity |
| mA | - | Mili Ampere |
| mA/cm ² | - | Curent Density Unit |
| Mg | - | Magnesium |
| mg | - | Mili gramme |
| MgO | - | Magnesium Oxide |
| Min | - | Minute |

| | | |
|-----------------|---|--|
| ml | - | Mili Litre |
| mm | - | Mili Meter |
| MOS | - | Metal Oxide Semiconductors |
| MOSFET | - | Metal Oxide Semiconductor Field-effect Transistors |
| mS | - | Mili Siemens |
| mS/cm | - | Conductivity Unit |
| mV | - | millivolt |
| N | - | Nitrogen |
| Na | - | Natrium |
| NaOH | - | Natrium Hydroxide |
| NH ₂ | - | Amine Functional Group |
| NH ₃ | - | Ammonia |
| NH ₄ | - | Ammonium |
| Ni | - | Nickel |
| nm | - | Nono Meter |
| NTR | - | Non-treated |
| O ₂ | - | Oxygen |
| OH | - | Hydroxyl Functional Group |
| OhBN | - | Organized Hexagonal BN |
| PAA | - | Polyacrylic Acid |
| PC 1 | - | Polycationic 1 |
| PC 2 | - | Polycationic 2 |
| PCB | - | Printed Circuit Board |
| PDADMAC | - | Poly(Diallyldimethylammonium Chloride) |

| | | |
|------------------|---|---------------------------------|
| PEG | - | Poly Ethylene Glycol |
| PEI | - | Polyethylene Imine |
| PVA | - | Poly Vinyl Alcohol |
| PZT | - | Lead Zirconate Titanate |
| R^2 | - | R Square |
| r-BN | - | Rhombohedral Boron Nitride |
| Rth | - | Temperature Resistance |
| SEM | - | Scanning Electron Microscopy |
| SiO ₂ | - | Silicon Dioxide |
| Si-OH | - | Silanol Functional Group |
| SnO ₂ | - | Stannum Oxide |
| t | - | Time |
| TC | - | Thermal Conductivity |
| TEOS | - | Tetraethyl Orthosilicate |
| Ti-6Al-4V | - | Titanium – Aluminium – Vanadium |
| TIM | - | Thermal Interface Material |
| TiO ₂ | - | Titanium Dioxide |
| TO | - | Transistor Outline |
| TR | - | Treated, |
| UPS | - | Uninterruptible Power Supply |
| V | - | Particle Velocity |
| V | - | Voltage |
| W | - | Waat |
| W | - | Weight |